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- (71) Applicant (for all designated States except US): **MEDITAB SPECIALITIES PVT. LTD** [IN/IN]; 12 Gunbow Street, Mumbai 400 001 (IN).
- (71) Applicant (for MW only): **WAIN, Christopher, Paul** [GB/GB]; A A Thornton & Co., 235 High Holborn, London WC1V 7LE (GB).
- (72) Inventors; and
- (75) Inventors/Applicants (for US only): **UTTARWAR, Sunil, Govindrao** [IN/IN]; Meditab Specialities Private Limited, D-22, MIDC Kurkumbh, Dist. Pune 413801, Maharashtra (IN). **GAWLI, Bhagwan, Narayan** [IN/IN]; Meditab Specialities Privat Limited, D-22, MIDC Kurkumbh, Dist. Pune 413801, Maharashtra (IN).
- (74) Agents: **WAIN, Christopher, Paul et al.**; A A Thornton & Co, 235 High Holbon, London WC1V 7LE (GB).
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(57) Abstract: A process of purifying citalopram, either in racemic or enantiomeric form, which process comprises (i) providing a crude mixture comprising citalopram, either in racemic or enantiomeric form, dissolved in a water immiscible organic solvent, and which mixture also includes one or more citalopram derivatives which are present as citalopram impurities; (ii) washing the crude mixture with at least one dilute aqueous solution of a polybasic acid, either in free form or as a partial alkali metal salt, so as to separate citalopram from citalopram impurities present in the crude mixture; and (iii) where required converting citalopram free base, separated from citalopram impurities further to step (ii), to a pharmaceutically acceptable salt.

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